Amendments to the Claims:

Please cancel claims 16 and 18.

Please amend claims 4-14, 17, and 20 as shown below.

This listing of claims replaces all prior versions and listings of claims in the application:

1. (original) A compound of the formula

$$^{8}RO$$
 $^{R^{6}}$
 $^{O}R^{7}$
 $^{O}R^{7}$
 $^{O}R^{5}$
 $^{O}R^{5}$

in which

is hydrogen, alkyl, aryl, heteroaryl, heterocyclyl, alkylcarbonyl, arylcarbonyl, heterocyclylcarbonyl, heteroarylcarbonyl, alkoxycarbonyl, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, alkylsulfonyl, arylsulfonyl, heterocyclylsulfonyl, heteroarylsulfonyl or a carbonyl-linked amino acid residue,

where R¹ apart from hydrogen may be substituted by 0, 1, 2, or 3 substitutents R¹⁻¹, where the substituents R¹⁻¹ are selected independently of one another from the group consisting of halogen, alkyl, trifluoromethyl,

trifluoromethoxy, nitro, cyano, amino, alkylamino, dialkylamino, cycloalkyl, aryl, heterocyclyl, hydroxy, alkoxy, and carboxyl,

R² is hydrogen or alkyl,

where alkyl may be substituted by 0,1,2, or 3 substituents R²⁻¹, where the substituents R²⁻¹ are selected independently of one another from the group consisting of halogen, amino, alkylamino and dialkylamino,

or

- R¹ and R² together with the nitrogen atom to which they are bonded form a heterocycle which may be substituted by 0, 1, or 2 substituents R¹⁻², where the substituents R¹⁻² are selected independently of one another from the group consisting of halogen, trifluoromethyl, amino, alkylamino, dialkylamino, cycloalkyl, aryl, heteroaryl, heterocyclyl, hydroxy, alkoxy, carboxyl, alkoxycarbonyl and aminocarbonyl,
- R³ is hydrogen, alkyl or the side group of an amino acid, in which alkyl may be substituted by 0,1,2, or 3 substituents R³⁻¹, where the substituents R³⁻¹ are selected independently of one another from the group consisting of trifluoromethyl, nitro, amino, alkylamino, dialkylamino, cycloalkyl, aryl, heteroaryl, heterocyclyl, hydroxy, alkoxy, carboxyl, alkoxycarbonyl, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, guanidino and amidino,

in which cycloalkyl, aryl, heteroaryl and heterocyclyl may be substituted by 0, 1 or 2 substituents R³⁻², where the substituents R³⁻² are selected independently of one another from the group consisting of halogen, alkyl, trifluoromethyl and amino,

and in which one or more free amino groups in the side group of the amino acid may be substituted by alkyl, alkenyl, alkynyl, cycloalkyl, aryl, heteroaryl, heterocyclyl, alkylcarbonyl, arylcarbonyl, heteroarylcarbonyl, heterocyclylcarbonyl, alkoxycarbonyl, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, arylaminocarbonyl, alkylsulfonyl, arylsulfonyl, heterocyclylsulfonyl or heteroarylsulfonyl,

- R³' is hydrogen, C₁-C₆-alkyl or C₃-C₈-cycloalkyl,
- R⁴ is hydrogen, C₁-C₆-alkyl or C₃-C₈-cycloalkyl,
- is alkyl, cycloalkyl, aryl, heteroaryl, heterocyclyl or a hydroxy function-linked amino acid residue, where R⁵ may be substituted by 0, 1, 2 or 3 substituents R⁵⁻¹, where the substituents R⁵⁻¹ are selected independently of one another from the group consisting of halogen, alkyl, trifluoromethyl, trifluoromethoxy, cyano, amino, alkylamino, dialkylamino, cycloalkyl, aryl, heteroaryl, heterocyclyl, hydroxy, alkoxy, carboxyl, alkoxycarbonyl, aminocarbonyl, alkylaminocarbonyl and dialkylaminocarbonyl,

in which alkylamino and dialkylamino may be substituted by 0, 1, or 2 substituents R⁵⁻², where the substituents R⁵⁻² are selected independently of one another from the group consisting of hydroxy, amino, alkoxy, alkylamino and dialkylamino,

- R⁶ is hydrogen, C₁-C₆-alkyl or C₃-C₈-cycloalkyl,
- R⁷ is hydrogen, C₁-C₆-alkyl, alkylcarbonyl or C₃-C₈-cycloalkyl,
- R^8 is hydrogen or C_1 - C_6 -alkyl,

and one of their salts, their solvates and the solvates of their salts.

2. (original) The compound as claimed in claim 1, characterized in that it corresponds to the formula

$$^{8}RO$$
 R^{6}
 $R^{1}R^{2}N$
 R^{3}
 R^{3}
 R^{4}
 OR^{7}
 OR^{7}
 OR^{5}

in which R¹ to R⁸ have the same meanings as in formula (I).

- 3. (original) The compound as claimed in claim 1 or 2, characterized in that
 - R¹ is hydrogen, alkyl or alkylcarbonyl,
 - R² is hydrogen,
 - R³ is alkyl or the side group of an amino acid, in which alkyl may be substituted by 0,1,2, or 3 substituents R³⁻¹, where the substituents R³⁻¹ are selected independently of one another from the group consisting of trifluoromethyl, nitro, amino, alkylamino, dialkylamino, cycloalkyl, aryl, heteroaryl, heterocyclyl, hydroxy, alkoxy, carboxyl, alkoxycarbonyl, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, guanidino and amidino,

in which cycloalkyl, aryl, heteroaryl and heterocyclyl may be substituted by 0, 1 or 2 substituents R³⁻², where the substituents R³⁻² are selected independently of one another from the group consisting of halogen, alkyl, trifluoromethyl and amino,

and in which one or more free amino groups in the side group of the amino acid may be substituted by alkyl,

- R^{3'} is hydrogen, C₁-C₆-alkyl or C₃-C₈-cycloalkyl,
- R⁴ is hydrogen, C₁-C₆-alkyl or C₃-C₈-cycloalkyl,
- is alkyl, cycloalkyl, aryl, heteroaryl or heterocyclyl, where R⁵ may be substituted by 0, 1, 2 or 3 substituents R⁵⁻¹, where the substituents R⁵⁻¹ are selected independently of one another from the group consisting of halogen, alkyl, trifluoromethyl, trifluoromethoxy, cyano, amino, alkylamino, dialkylamino, cycloalkyl, aryl, heteroaryl, heterocyclyl, hydroxy, alkoxy, carboxyl, alkoxycarbonyl, aminocarbonyl, alkylaminocarbonyl and dialkylaminocarbonyl,

in which alkylamino and dialkylamino may be substituted by 0, 1, or 2 substituents R⁵⁻², where the substituents R⁵⁻² are selected independently of one another from the group consisting of hydroxy, amino, alkoxy, alkylamino and dialkylamino,

- R⁶ is hydrogen,
- R⁷ is hydrogen, C₁-C₆-alkyl, alkylcarbonyl or C₃-C₈-cycloalkyl,

and

 R^8 is hydrogen. (amended) The compound as claimed in any of claims 1 to 3 claim 3, 4. characterized in that R^1 is hydrogen, R^2 is hydrogen, \mathbb{R}^3 is aminocarbonylmethyl, 3-aminoprop-1-yl, 2-hydroxy-3-aminoprop-1-yl, 1hydroxy-3-aminoprop-1-yl, 3-guanidinoprop-1-yl, 2-aminocarbonylethyl, 2hydroxycarbonylethyl, 4-aminobut-1-yl, hydroxymethyl, 2-hydroxyethyl, 2aminoethyl, 4-amino-3-hydroxybut-1yl or (1-piperidin-3-yl)methyl, R^{3} is hydrogen, R^4 is hydrogen, methyl, ethyl, isopropyl or cyclopropyl, is alkyl or C₃-C₆-cycloalkyl, where R⁵ may be substituted by 0, 1, 2 or 3 R^5 substituents R⁵⁻¹, where the substituents R⁵⁻¹ are selected independently of one another from the group consisting of alkyl, amino, alkylamino, dialkylamino, cycloalkyl, hydroxy, alkoxy, carboxyl, alkoxycarbonyl, aminocarbonyl, alkylaminocarbonyl and dialkylaminocarbonyl, in which alkylamino and dialkylamino may be substituted by 0, 1, or 2 substituents R⁵⁻², where the substituents R⁵⁻² are selected independently of one another from the group consisting of hydroxyl and amino,

 R^6

 R^7

is hydrogen,

is hydrogen,

and	
R ⁸	is hydrogen.
(amer	nded) The compound as claimed in any of claims 1 to <u>claim</u> 4, characterized t
R ¹	is hydrogen,
R ²	is hydrogen,
R^3	is 3-aminoprop-1-yl or 2-hydroxy-3-aminoprop-1-yl,
R³'	is hydrogen,
R ⁴	is hydrogen or methyl,
R ⁵	is C_1 - C_4 -alkyl where alkyl may be substituted by 0, 1 or 2 substituents independently of one another selected from the group consisting of amino hydroxyl and carboxyl,
R^6	is hydrogen,
R ⁷	is hydrogen,
and	
R ⁸	is hydrogen.

5.

- 6. (amended) The compound as claimed in any of claims 1 to 3 claim 1 or 2, characterized in that
 - R¹ is hydrogen.
- 7. (amended) The compound as claimed in any of claims 1, 2 and 6 claim 1 or 2, characterized in that R² is hydrogen.
- 8. (amended) The compound as claimed in any of claims 1 to 4, 6 and 7 claim 1 or 2, characterized in that R³ is 3-aminoprop-1-yl or 2-hydroxy-3-aminoprop-1-yl.
- 9. (amended) The compound as claimed in any of claims 1 to 3 or 6 to 8 <u>claim 1 or</u> <u>2</u>, characterized in that R^{3'} is hydrogen.
- (amended) The compound as claimed in any of claims 1 to 4 or 6 to 9 claim 1 or
 characterized in that R⁴ is hydrogen or methyl.
- 11. (amended) The compound as claimed in any of claims 1 to 4 or 6 to 10 claim 1 or 2, characterized in tht R⁵ is C₁-C₄-alkyl, where alkyl may be substituted by 0, 1 or 2 substituents independently of one another selected from the group consisting of amino, hydroxyl and carboxyl.
- 12. (amended) The compound as claimed in any of claims 1, 2, 6 to 11 <u>claim 1 or 2</u>, characterized in that R⁶ is hydrogen.
- 13. (amended) The compound as claimed in any of claims 1 to 3 or 6 to 12 claim 1 or 2, characterized in that R⁷ is hydrogen.
- 14. (amended) The compound as claimed in any of claims 1, 2, 6 to 13 <u>claim 1 or 2</u>, characterized in that R⁸ is hydrogen.

15. (original) A process for preparing a compound of the formula (I) as claimed in claim 1, characterized in that a compound of the formula

$$^{8}RO$$
 R^{6}
 $R^{1}R^{2}N$
 R^{3}
 R^{3}
 R^{4}
 R^{4}
 R^{4}
 R^{4}
 R^{5}
 R^{4}
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 R^{5}
 R^{5}
 R^{4}
 R^{5}
 R^{5}

in which

R¹ to R⁴ and R⁶ to R⁸ have the meaning indicated in claim 1,

is reacted with a compound of the formula

$$HO - R^5$$
 (III),

in which

R⁵ has the meaning indicated in claim 1.

- 16. (canceled)
- 17. (amended) A medicament comprising at least one compound as claimed in any of claims 1 to 14 claim 1 or 2 in combination with at least one pharmaceutically suitable, pharmaceutically acceptable carrier or other excipients.
- 18. (canceled)

- 19. (original) A medicament as claimed in claim 17 for the treatment and/or prophylaxis of bacterial infections.
- 20. (amended) A method for controlling bacterial infections in humans and animals by administration of an antibacterially effective amount of at least one compound as claimed in any of claims 1 to 14 claim 1 or 2.